

DT 24-APR-2001 (first entry)
 XX
 DE Human RECAP polypeptide, SEQ ID NO: 21.
 XX
 KW Human; RECAP; receptors and associated proteins; cerebroprotective;
 KW nootropic; neuroprotective; anticonvulsant; antiparkinsonian; anti-HIV;
 KW antidiabetic; immunostimulant; immunomodulator; antiinflammatory;
 KW antithyroid; immunosuppressive; nephrotropic; antigout; thyromimetic;
 KW cytostatic; antibacterial; virucide; fungicide; protozoacide;
 KW antiarteriosclerotic; hepatotropic; gene therapy; infection; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO200107612-A2.
 XX
 PD 01-FEB-2001.
 XX
 PF 21-JUL-2000; 2000WO-US020035.
 XX
 PR 21-JUL-1999; 99US-0145232P.
 PR 07-OCT-1999; 99US-0158578P.
 PR 12-NOV-1999; 99US-0165192P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Au-Young J, Bandman O, Tang YT, Yue H, Azimzai Y, Burford N;
 PI Baughn MR, Lu DAM, Hillman JL, Patterson C, Lal P;
 XX
 DR WPI; 2001-168554/17.
 DR N-PSDB; AAF58615.
 XX
 PT Novel receptors and associated proteins for diagnosis and treatment of
 PT neurological disorders, immunological disorders including autoimmune/
 PT inflammatory disorders and cell proliferative disorders such as cancer.
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 PS Disclosure; Page 111-112; 128pp; English.
 XX
 CC The present sequence is a human RECAP (receptors and associated proteins)
 CC polypeptide. RECAP polynucleotides and polypeptides are useful in the
 CC diagnosis, treatment and prevention of neurological disorders such as
 CC stroke, Alzheimer's disease, Pick's disease, Huntington's disease,
 CC dementia, Parkinson's disease, Downs's syndrome, amyotrophic lateral
 CC sclerosis, multiple sclerosis, bacterial and viral meningitis, CJD
 CC (Creutzfeldt-Jakob disease), GSS (Gerstmann -Straussler-Scheinker
 CC syndrome); immunological disorders, including autoimmune/inflammatory
 CC disorders such as AIDS, DiGeorge's syndrome, severe combined
 CC immunodeficiency disease (SCID), Chediak-Higashi syndrome, Cushing's
 CC disease, Addison's disease, autoimmune thyroiditis, Crohn's disease,
 CC diabetes mellitus, Good pasture's syndrome, gout, Grave's diseases,
 CC Hashimoto's thyroiditis, Sjogren's syndrome, Werner's syndrome, viral,
 CC bacterial, fungal, parasitic, protozoal, and helminthic infections; and
 CC cell proliferation disorders such as arteriosclerosis, atherosclerosis,
 CC cirrhosis, hepatitis and cancer
 XX
 SQ Sequence 357 AA;

Query Match 100.0%; Score 1865; DB 4; Length 357;
 Best Local Similarity 100.0%; Pred. No. 2.7e-203;
 Matches 357; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MATTVPDGCNRGLKSKYYRLCDKAEAWGIVLETVATAGVVTSAFMLTLPILVCKVQDSN 60

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Db      1 MATTVPDGCNRGLKSKYYRLCDKAEAWGIVLETVATAGVVTSAFMLTLPILVCKVQDSN 60
Qy      61 RRKMLPTQFLFLGLGVLGIFGLTFAFIIGLDGSTGPTREFFLFGILFSICFSCLLAHAVSLT 120
Db      61 RRKMLPTQFLFLGLGVLGIFGLTFAFIIGLDGSTGPTREFFLFGILFSICFSCLLAHAVSLT 120
Qy      121 KLVRGRKPLSLLVILGLAVGFSLVQDVIAIEYIVLTMNRTNVNVFSELSAPRRNEDFVLL 180
Db      121 KLVRGRKPLSLLVILGLAVGFSLVQDVIAIEYIVLTMNRTNVNVFSELSAPRRNEDFVLL 180
Qy      181 LTYVFLMALTFMLSSFTFCGSFTGWKRHGAHIYLTMLLSIAIWVAWITLLMLPDFDRRW 240
Db      181 LTYVFLMALTFMLSSFTFCGSFTGWKRHGAHIYLTMLLSIAIWVAWITLLMLPDFDRRW 240
Qy      241 DDTILSSALAANGWVFLLAYVSPEFWLLTKQRNPMDYPVEDAFCKPQLVKKSYGVENRAY 300
Db      241 DDTILSSALAANGWVFLLAYVSPEFWLLTKQRNPMDYPVEDAFCKPQLVKKSYGVENRAY 300
Qy      301 SQEEITQGFEEETGDTLYAPYSTHFQLQNQPPQKEFSIPRAHAWPSPYKDYEVKKEGS 357
Db      301 SQEEITQGFEEETGDTLYAPYSTHFQLQNQPPQKEFSIPRAHAWPSPYKDYEVKKEGS 357

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RESULT 2

AAB93311

ID AAB93311 standard; protein; 357 AA.

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AC AAB93311;

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DT 26-JUN-2001 (first entry)

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DE Human protein sequence SEQ ID NO:12389.

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KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.

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OS Homo sapiens.

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PN EP1074617-A2.

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PD 07-FEB-2001.

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PF 28-JUL-2000; 2000EP-00116126.

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PR 29-JUL-1999; 99JP-00248036.

PR 27-AUG-1999; 99JP-00300253.

PR 11-JAN-2000; 2000JP-00118776.

PR 02-MAY-2000; 2000JP-00183767.

PR 09-JUN-2000; 2000JP-00241899.

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PA (HELI-) HELIX RES INST.

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PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX

DR WPI; 2001-318749/34.

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PT Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs.